

IN THE CLAIMS

This listing of claims will replace all prior versions and listings of claims in the application.

Claims 1-8 are pending in the application. Please amend claims 1, 3, 5 and 7 as follows:

1. (Currently Amended) An active implantable medical device comprising:
means for sensing ventricular and atrial cardiac activity,
analyzing means for discriminating a ventricular extrasystole from said sensed ventricular cardiac activity, and for detecting in the sensed cardiac activity an occurrence of a cardiac cycle,
measuring means for delivering an intracardiac impedance signal measured between two cavities of the heart correlated to the instantaneous blood flow, and
discriminating means for detecting variations in the delivered impedance signal measured between two cavities of the heart, determining, in response to a detected variation, a presence or an absence of a significant mechanical activity of the myocardium consecutive to a discriminated ventricular extrasystole, and determining a noxious character of the ventricular extrasystole in response to a determined absence of significant mechanical activity and a non noxious character of the ventricular extrasystole in response to a determined presence of significant mechanical activity, wherein the discriminating means further comprises means for evaluating an increase in the intracardiac impedance measured between two cavities of the heart after the occurrence of the detected ventricular extrasystole, comparing said increased intracardiac impedance measured between two cavities of the heart to a predetermined threshold, and determining a presence of a significant mechanical activity of the myocardium in response to said increase crossing said threshold.

2. (Original) The device of claim 1, wherein the device further comprises an operating parameter associated with controlling a cardiac rhythm and further comprises means for modifying said operating parameter for a cardiac cycle following the detected ventricular extrasystole in a first manner in response to a determined noxious character of the detected ventricular extrasystole and in a second manner for a determined non noxious character of the detected ventricular extrasystole.
3. (Currently Amended) The device of ~~the~~ claim 2, wherein said device further comprises means for calculating a ventricular escape interval and said operating parameter is said calculated ventricular escape interval.
4. (Cancelled)
5. (Currently Amended) The device of claim [4] 1, wherein said measuring means further comprises means for measuring an initial intracardiac impedance measured between two cavities of the heart and measured at the time of the occurrence of the ventricular extrasystole, and said predetermined threshold corresponds to the measured initial intracardiac impedance measured between two cavities of the heart increased by a multiplier greater than one.
6. (Original) The device of claim 1, further comprising counting means for counting a number of determined noxious extrasystoles and a number of determined non noxious extrasystoles.
7. (Currently Amended) The device of ~~the~~ claim 6, wherein said counting means further comprises means for determining a parameter representative of a relative proportion of counted noxious and non noxious extrasystoles.
8. (Previously Presented) The device of claim 5, wherein said multiplier is set to 1.5.